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(64) Water reducible coating compositions.

(57) An aqueous dispersion of a tertiary amine terminated polyether and a carboxylic acid functional polymer. The tertiary amine terminated polyether is the product of the reaction of the epoxy group of an epoxy resin with the hydroxy group of a hydroxy functional tertiary amine and the carboxylic acid functional polymer is present in an amount sufficient to render the combination water dispersible.

A process for producing the aqueous dispersion is also described, together with coating compositions based on the dispersion. These coating compositions are particularly useful in the lining of metal containers for food and beverages.

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WATER REDUCIBLE COATING COMPOSITIONS

This invention relates to an aqueous dispersion of a tertiary amine terminated polyether and a carboxylic acid functional polymer, a process for producing the same and coating compositions based on the aqueous dispersion of polymer. The coating compositions of this invention are inter alia useful for lining metal containers for food, beer and other beverages.

Epoxy resins are generally regarded as providing an excellent combination of properties; offering good adhesion, flexibility and resistance to solvent and chemical attack.

There is an increasing demand, worldwide, for low V.O.C. water reducible coating compositions to replace solvent based systems. [V.O.C. is the weight of organic solvent per unit volume of solid coating, expressed in grams per litre]. Epoxy resins are hydrophobic in nature. Consequently, many attempts have been made to modify epoxy resins to make them water dispersible without loss of their desirable properties. These approaches have met with varying degrees of success.

For example, EP 6334, EP 116225 and related patents teach the grafting of an epoxy resin to a hydrophilic acrylic polymer containing carboxyl groups, by reaction between the epoxide groups of the epoxy resin and the carboxylic acid groups of the acrylic polymer. Thus the bonds linking the hydrophilic and hydrophobic components of the copolymer are ester bonds. This approach has the disadvantage that, in a water based coating composition, the ester bonds tend to hydrolyse, resulting in poor storage stability.

British Patent Nos. 1,585,486 and 1,585,487 teach the approach of growing a hydrophilic acrylic

polymer containing carboxyl groups by free radical addition polymerisation, in the presence of a modified epoxy resin. The key to this approach is in the use of relatively high levels of a free radical initiator

5 which has high hydrogen abstraction power (principally benzoyl peroxide); this leads to the grafting of the acrylic polymer to the epoxy resin by chain transfer.

This approach has the disadvantage that the level of grafting is relatively low. Furthermore, the water

10 based coating composition made by this approach has a relatively high viscosity for a given solids.

Yet a different approach is taught by

U.S. 4,247,439, WO 83/03613 and related patents. These patents teach the reaction of the residual epoxide

15 groups of an epoxy resin with a tertiary amine, in the presence of water, to form a quaternary amine hydroxide terminated polyether. The product of this reaction is

dispersed in water by incorporating a blend of carboxylic acid functional polymers. This approach has

20 the great disadvantage that it is cumbersome, and hence expensive, in that there are a relatively large number of intermediate products that go to make up the final product.

According to the present invention there is
25 provided an aqueous dispersion of a tertiary amine terminated polyether and a carboxylic acid functional polymer wherein the tertiary amine terminated polyether is the product of the reaction of the epoxy group of an epoxy resin with the hydroxy group of a hydroxy
30 functional tertiary amine and the carboxylic acid functional polymer is present in an amount sufficient to render the combination water dispersible.

The invention also provides a process for producing this aqueous dispersion which comprises:

35 a) reacting the epoxide groups of an epoxy resin with a hydroxy functional tertiary amine to

form a tertiary amine terminated polyether, and

- b) dispersing the tertiary amine terminated polyether in water with the aid of a carboxylic acid functional polymer in an amount
5 sufficient to render the combination dispersible.

There is further provided a coating composition comprising a stable dispersion in water of:

- 1) a tertiary amine terminated polyether which is in ionic association with
10 ii) a polymer derived, at least in part, from carboxylic acid functional monomers, and
iii) a crosslinking agent.

It will be appreciated that, in practice, some unassociated polymer of type i) and/or type ii) may
15 also be present in the composition.

An ionically associated polymer complex is thought to result from an acid base relationship between the amine terminated polyether and the carboxylic acid containing polymer. The amine
20 terminated polyether is formed by reacting the 1,2 epoxide group of an epoxy resin with the hydroxy group of a hydroxy functional tertiary amine (e.g. N,N dimethyl ethanolamine). Optionally the coating may also contain zinc oxide or other pigmentation, depending on
25 the final application.

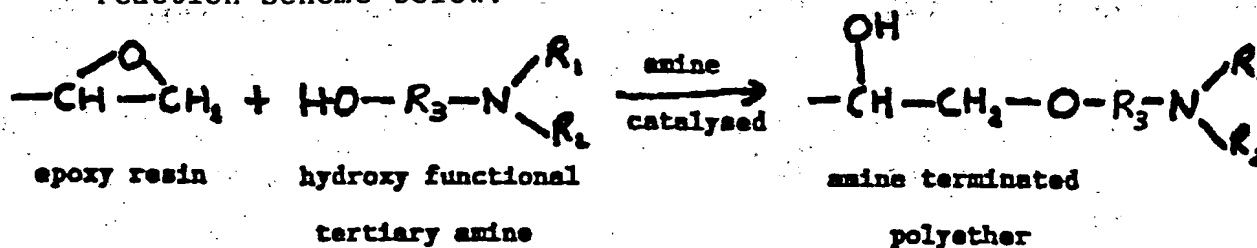
An advantage of this product is that it allows considerable flexibility in formulating coating compositions with desired solids viscosity relationships. Typical examples would be: 45% solids
30 at a viscosity of 26 sec. Ford No.4 Cup at 25°C, and 17-1/2% solids at a viscosity of 21 sec. Ford No. 4 Cup.

In the simplest approach to making the amine terminated polyether used in the present invention, a
35 low molecular weight epoxy resin (e.g. DER343) is reacted with Bisphenol A in a suitable organic solvent.

The hydroxy functional tertiary amine is added to the resultant epoxy resin and the reaction mixture is held at ca 100°C for about 1 hour. Examples of suitable hydroxy functional tertiary amines are

N,N dimethylethanolamine (DMEA), N,N diethylethanolamine (DEEA), 2-dimethylamino-2-methyl-propanol (DMAMP) and triethanolamine.

During the course of the reaction, the tertiary amine group of the hydroxy functional amine catalyses the reaction of the hydroxy group with the residual epoxide groups of the epoxy resin, according to the reaction scheme below:

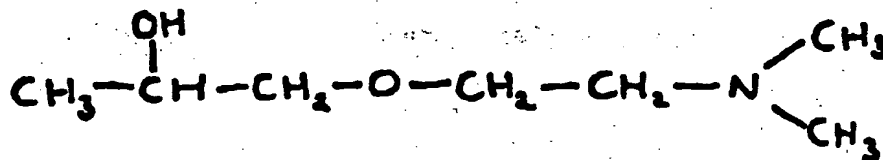


Where, in the case of DMEA, R₁ and R₂ are methyl groups and R₃ is -CH₂-CH₂-. It must be borne in mind that the presence of a hydroxy functional solvent (e.g. butanol or butyl cellosolve) will reduce the yield of the tertiary amine terminating groups by competing with the hydroxyl groups of the tertiary amine. If significant amounts of water are present, the undesirable quaternary amine hydroxide will be formed. Thus preferred results are achieved when the reaction is carried out under anhydrous conditions.

In order to confirm the reaction products stated above, we carried out experiments in which, in order to facilitate analysis, we maximised the number of 1,2 epoxide groups available for reaction.

EXPERIMENT 1REACTION OF DMEA WITH EPOXIDE GROUPS

5 Dry propylene oxide, 146.6g, and dry DMEA 224.9g (water content less than 0.05%) were mixed together in a round-bottomed flask fitted with a thermometer, a stirrer and a reflux condenser. The reaction mixture was heated to 40°C whereupon it exothermed to 90°C before the temperature began to fall. Excess reactants were distilled off using a rotary evaporator. Analysis of the product by NMR and GPC showed that, despite some polymerisation, the reaction product was predominantly that shown below:

EXPERIMENT 2

15 REACTION OF DRY DMEA WITH THE
DIGLYCIDYL ETHER OF BISPHENOL A

20 Dry DER331 11g, and dry DMEA 211g were mixed together in a 500 ml round-bottomed flask fitted with a condenser, a thermometer, a stirrer and temperature controller, heated to 105°C and held at that temperature for 1 hour. Excess DMEA was removed from the reaction mixture using a rotary evaporator. Analysis by NMR showed that a considerable amount of the product obtained was due to the reaction of the hydroxyl group of DMEA with the epoxide group of DER331. Further evidence for the tertiary amine terminated product was that it was soluble in water only at low pH.

EXPERIMENT 3REACTION OF DMEA WITH THE DIGLYCIDYL ETHER
OF BISPHENOL A IN THE PRESENCE OF WATER

DER331 2.2g, DMEA 10.7g, water 208g, and
5 dimethyl formamide 283g were mixed together in a round-
bottomed flask fitted with a reflux condenser, a
stirrer and a thermometer and heated to reflux (106°C).
The mixture was maintained at reflux for 6 hours. The
product was purified using a rotary evaporator.

10 Analysis of the product was consistent with it being
quaternary amine salt. Furthermore, the product was
water soluble under neutral, acidic and basic
conditions.

In the product of this invention the tertiary
15 amine terminated polyether is dispersed in water with
the aid of a carboxylic acid functional polymer. The
number of carboxylic acid groups in the carboxylic acid
functional polymer in a given amount of product needs
to be sufficient to form an effective ionic association
20 with the amine terminated polyether and for a stable
dispersion in water to be formed. If the carboxylic
acid functional polymer is a copolymer of an
ethylenically unsaturated acid, such as acrylic acid or
methacrylic acid, and hydrophilic co-monomers such as
25 2 hydroxy propyl methacrylate, 2 hydroxy ethyl acrylate,
or acrylamide, then the number of acid groups on the
carboxylic acid functional polymer in a given amount
of product may be less than the number of tertiary
amine groups on the polyether in that same amount of
30 product. If, however, the carboxylic acid functional
polymer is a copolymer of an ethylenically unsaturated
acid and more hydrophobic monomers such as styrene or
butyl acrylate, then a larger proportion of acid groups

is required than if hydrophilic co-monomers are used.

Our experience with this type of system reveals that the final viscosity of the coating composition, at a given amount of solids, is governed by:

- 5 i) The spacing of the carboxylic acid groups in the carboxylic acid functional polymer. The more the acid groups are spaced by other monomers, at a given total number of carboxylic groups, the lower the viscosity.
- 10 ii) The VOC; the lower the VOC, the lower the viscosity.
- 15 iii) The degree of neutralisation of the carboxylic acid groups; the lower the degree of neutralisation, the lower the viscosity. It is always necessary, however, to ensure a suitable degree of neutralisation to provide a stable dispersion.
- 20 iv) The ratio of carboxylic acid functional polymer to the tertiary amine terminated polyether. The higher the ratio of carboxylic acid functional polymer, the higher the viscosity.
- v) The molecular weight of the carboxylic acid functional polymer; the higher the molecular weight, the higher the viscosity.

As mentioned above, the role of the solvent can be important in determining the properties of the final coating composition. If the degree of ionic association between the polyether and the polycarboxylic acid functional polymer is too great, the polymer blend will gell and it will no longer be possible to disperse it in water. The degree of ionic association is controlled by controlling the amine value of the polyether; more specifically by minimising the number of molecules which have tertiary amine groups on both ends. This can be achieved by, in the first instance, minimising the number of modified epoxy resin molecules with 1,2 epoxide groups on each end, and in the second instance, by reacting some of these

groups with hydroxy functional solvents, such as butyl cellosolve or butanol. Obviously, if the amine termination reaction is carried out in the presence of an hydroxy functional solvent, the extent of amine termination would be reduced because the tertiary amine would be expected to catalyse the reaction of the solvent hydroxyl groups with the epoxide groups of the epoxy resin.

EXAMPLE 1

CARBOXYLIC ACID FUNCTIONAL POLYMER
FOR A HIGH SOLIDS COATING

		<u>Wt. (g)</u>
a a	Acrylic acid	584
a b	Styrene	843
15 a c	Benzoyl peroxide	7.0
b b	Butyl cellosolve	1414
b c	Ethyl alcohol	1106
c c	t-butyl perbenzoate	5.4
c d	Toluene	18.4
20 d d	t-butyl perbenzoate	5.4
d e	Toluene	18.4

1. aa, ab and ac were added to a 5-litre five-necked, flange-topped round-bottomed flask fitted with a condenser, a stirrer and a thermometer, and stirred until ac had dissolved.
2. bb and bc were then added and the mixture heated to reflux temperature (ca 96°C) and held at this temperature over 1 hour.

3. cc and cd were then premixed, added to the flask and held at reflux for 30 minutes.
4. dd and de were premixed, added to the flask and held at reflux temperature for 1 hour.

5

EXAMPLE 2HIGH SOLIDS, LOW VISCOSITY COATING COMPOSITION

			<u>Wt. (g)</u>
	a a	Epikote 8290 (Product of Shell Chemical Co.)	519
10	a b	Bisphenol A	470
	a c	Butyl cellosolve	285
	b b	DMEA	53
	b c	Butyl cellosolve	107
15	c c	Polyelectrolyte solution (Example 1)	1182
	d d	DMEA	70
	e e	Water	1888
	f f	Cymel 303 (Product of American Cyanamid Co.)	427

- 20 1. aa, ab and ac were added to a 5-litre five-necked, flange-topped, round-bottomed flask fitted with a condenser, a stirrer and a thermometer, and mixed well. The vessel contents were heated to 170°C over 1 hour; the heating was turned off and the exotherm was allowed

5 to take the temperature to reflux (195-200°C). The reaction was allowed to proceed until a viscosity of W at 25°C was obtained (diluted to 52% solids in butyl cellosolve), (total cook time was 1.5 hours), after which the vessel contents were cooled to 130°C over a period of 45 minutes.

2. bb and bc were then premixed and added slowly to the flask with stirring. The vessel contents were cooled to 100°C and held at that temperature for 10 1 hour. About 150g of solvent was removed by distillation under an air stream to remove excess amine.

15 3. cc was then added to the vessel and mixed in well. A further 736g of solvent was distilled off using an air stream.

4. dd was added and mixed in well.

5. ee was added and mixed in well.

6. ff was added and mixed in well for 20 minutes.

20 The finished coating composition had 45% solids, a VOC of 166g/litre and a viscosity of 26 sec. Ford 4 Cup @ 25°C. A sample of this coating composition was drawn down on aluminium foil and cured for 10 minutes at 200°C. The coating exhibited excellent film properties.

EXAMPLE 3

WATER BASED CARBOXYLIC ACID FUNCTIONAL POLYMER

			<u>Wt. (g)</u>
	a a	Acrylic acid	477
5	a b	Acrylamide	115
	a c	Water	3280
	b b	Potassium persulphate	6
	b c	Water	120

1. aa, ab and ac were added to a 5-litre five-necked,
10 flange-topped, round-bottomed flask fitted with a
condenser, a stirrer and a thermometer, and heated to
reflux (100°C). The heating was then turned off.
2. bb and bc were premixed and added to the flask
over 2 hours; the heating was then turned on and
15 the vessel contents were held at reflux for 1 hour.

EXAMPLE 4

A LOW SOLIDS AND HIGH VISCOSITY
COATING COMPOSITION BASED ON CARBOXYLIC ACID
FUNCTIONAL POLYMER OF EXAMPLE 3

		<u>Wt. (g)</u>
5		
	a a	Epikote 8920
	a b	Bisphenol A
	a c	Butyl cellosolve
	b b	DMEA
10	b c	Butyl cellosolve
	c c	Polyelectrolyte solution (Example 3)
	d d	DMEA
	e e	Water
15	f f	Cymel 303

The method of preparation was as for Example 2 except that no solvent was distilled off in Step 3.

The resulting coating composition had a viscosity of 78 sec. Ford Cup No.4 at 25°C when adjusted to 17% solids, with water. A sample of this coating composition was drawn down on aluminium foil and cured for 10 minutes at 200°C. The coating exhibited excellent film properties.

EXAMPLE 5

CARBOXYLIC ACID FUNCTIONAL POLYMER WITH
RELATIVELY LOW CONCENTRATION OF ACID GROUPS

			<u>Wt. (g)</u>
5	a a	Methoxy propoxy propanol	400
	a b	Deionised water	4.4
	a c	Methyl Ethyl Ketone	45
	a d	Paxwax 6364LA (National Wax Co.)	40
	b b	Butyl Acrylate	680
10	b c	Acrylic acid	65
	b d	Hydroxy propyl methacrylate	210
	c c	Benzoyl peroxide	4.5
	c d	Methyl ethyl ketone	45
	d d	Tertiary butyl perbenzoate	4
15	d e	Butyl cellosolve	40
	d g	Water	5
	e e	Tertiary butyl perbenzoate	4
	e f	Butyl cellosolve	50
	e g	Water	5
20	h h	N,N-dimethylethanolamine	40
	i i	Water	1385
	j j	Butyl cellosolve	223

1. aa, ab, ac and ad were added to a 5-litre five-necked, flange-topped, round-bottomed

flask fitted with a condenser, a stirrer, two dropping funnels and a thermometer-temperature controller, and heated to 100°C under a nitrogen blanket.

- 5 2. Add bb, bc and bd and cc and cd simultaneously over 2 hours while maintaining the temperature at 110°C.
3. Add dd, de, df and hold at 100°C for 1 hour.
4. Add ee, ef and eg and hold at 100°C for 1 hour.
- 10 5. Add hh and mix well.
6. Add ii and mix well. Distil off 1840g of solvent at temperatures between 100°C and 120°C.
7. Add jj and mix well.

EXAMPLE 6

15 COATING COMPOSITION BASED ON EXAMPLE 5

			<u>Wt. (g)</u>
	a a	DER343 (Dow Chemical Co.)	341
	a b	Bisphenol A	293
	a c	Butyl cellosolve	177
20	b b	N,N-dimethylethanolamine	34
	b c	Butyl cellosolve	34
	c c	Product of Example 5	707
	d d	Water	1600
	e e	Cymel 303 (Cyanamid Co.)	120
25	f f	Water	207
	g g	2% Phosphoric acid aqueous solution neutralized with N,N-dimethylethanolamine (pH = 7.0)	82

1. aa, ab and ac were added to a 5-litre five-necked, flange-topped, round-bottomed flask fitted with a condenser, a stirrer and a thermometer-temperature controller, and mixed well. The vessel contents were heated to 170°C, the heating was turned off and the exotherm was allowed to take the temperature to reflux (195-200°C). Allow reaction to cool to 170°C.
2. Cool to 130°C, add bb and bc; maintaining the temperature at 105°C for 1 hour. Distil off 34g of solvent.
3. Add cc and mix well at 70°C over 30 minutes.
4. Add dd slowly with good mixing. Cool to 50°C.
5. Add ee and mix well.
6. Add to adjust viscosity; mix well.
7. Add gg and mix until any precipitate redisperses.

The resulting coating composition had a viscosity of 50 sec. Ford Cup No. 4 at 25°C at 38% solids.

20

EXAMPLE 7

PREPARATION OF AMINE TERMINATED POLYETHER FOR USE IN EXAMPLES 8 AND 9

		<u>Wt. (g.)</u>
a a	DER343	681
25 a b	Bisphenol A	586
a c	Butyl cellosolve	355
b b	DMEA	67
b c	Butyl cellosolve	67

1. aa, ab and ac were mixed in a 2-litre flask fitted with a condenser, a thermometer-temperature

30

controller, and a stirrer, heated to 170°C and the heating turned off. The exotherm was allowed to take the temperature to 200°C. When the temperature decreased to 170°C, the reaction was cooled to 130°C.

5

2. bb and bc were premixed and added to the reaction mixture. The temperature was maintained at 105°C for 1 hour after which 67g of solvent was distilled off.

10

The following two examples demonstrate a method of preparing the carboxylic acid functional polymer in the presence of the amine terminated polyether.

EXAMPLE 8

USING LOW ACID VALUE CARBOXYLIC ACID FUNCTIONAL POLYMER

15

		<u>Wt. (g)</u>
	a a	Amine terminated polyether (Example 7)
	a b	Butanol
20	a c	Paxwax 6364LA
	b b	Butyl acrylate
	b c	Styrene
	b d	Acrylic Acid
	b e	Hydroxy propyl methacrylate
25	c c	Benzoyl peroxide
	c e	Toluene
	d d	Tertiary butyl perbenzoate
	d e	Butanol

(EXAMPLE 8 Cont 'd...)

		<u>Wt. (g)</u>
e e	Tertiary butyl perbenzoate	6
e f	Butanol	20
5 f f	Tertiary butyl perbenzoate	6
f g	Butanol	20
g g	Dimethylethanolamine	136
g h	Water	400
h h	Water	4002
10 i i	Cymel 303	212
j j	Water	4600

1. aa, ab and ac were mixed in a 5-litre flask fitted with a thermometer-temperature controller, a condenser, a stirrer and 2 droppers. The mixture was heated to 100°C.
- 15 2. bb, bc, bd and be; cc and ce were premixed, and slowly added into the flask over 2 hours at controlled temperature of 100°C. The temperature was maintained at 100°C for 1 hour.
- 20 3. dd and de were premixed and added and reaction was maintained at 110°C over 15 mins.
4. ee and ef were premixed and added, and the reaction was kept at 120°C over 1 hour.
- 25 5. ff and fg were premixed and added, cooked at 120°C over 6 hours and then cooled to 90°C.

Dispersion procedure:

6. gg was added with constant stirring.
7. hh was slowly added while mixing well.

8. ii was then added with stirring.

9. jj was added with stirring.

The above coating composition was diluted to 17.5% solids and was found to have a viscosity of 16 sec. Ford Cup No.4 at 25°C. The coating was sprayed into 2-piece aluminium cans using a Nordson laboratory spray machine and exhibited excellent coverage at both ca 100 mg and ca 180 mg per can.

EXAMPLE 9

10

USING HIGH ACID VALUE CARBOXYLIC ACID FUNCTIONAL POLYMER

			<u>Wt. (g)</u>
	a a	Amine terminated polyether (Example 7)	659
15	a b	Methoxy propoxy propanol	400
	a c	Paxwax 6364LA	20
	b b	Acrylic acid	169
	b c	Styrene	242
	c c	Benzoyl peroxide	4
20	c d	Toluene	40
	d d	Tertiary butyl perbenzoate	2
	d e	Methoxy propoxy propanol	20
	e e	Tertiary butyl perbenzoate	2
	e f	Methoxy propoxy propanol	20

1. aa, ab and ac were mixed in a 2-litre flask fitted with a thermometer-temperature controller, a condenser, a stirrer and 2 dropping funnels. The mixture was heated to 100°C.
- 5 2. bb and bc were premixed, and cc and cd were premixed. The two premixes were separately added over 1 hour 15 mins. at 100°C. The temperature, then, was increased to 120°C and maintained at that temperature over 45 mins.
- 10 3. dd and de were premixed and added. The reaction was maintained for 1 hour 15 mins. during which time the exotherm was allowed to take the temperature to 150°C.
- 15 4. ee and ef were premixed and added into the flask. The reaction was maintained at 120°C for 1 hour 15 mins. 140g of organic solvent was distilled to eliminate residual monomer.

The reaction product was dispersed in water in the same way as for Example 8.

- 20 The resulting coating composition had a viscosity of 40 sec. Ford Cup No. 4 at 25°C at 29.4% solids.

The following four Examples demonstrate the use of hydroxy functional tertiary amines other than DMEA.

25

EXAMPLE 10

PREPARATION OF COATING COMPOSITION BASED ON AMINE TERMINATED POLYETHER USING N,N-DIETHYLETHANOLAMINE (DEEA)

		<u>Wt. (g)</u>
30	a a	DER343
		213
	a b	Bisphenol A
		182
	a c	Butyl cellosolve
		111

(Example 10 cont'd...)

		<u>Wt. (g)</u>
b b	DEEA	27.6
b c	Butyl cellosolve	27.6
5 c c	Carboxylic acid functional polymer (Example 5)	468
d d	Water	777

1. aa, ab and ac were mixed in a 2-litre flask fitted with a thermometer-temperature controller, a condenser, a stirrer and heated to 170°C, at which temperature the heating was turned off. The exotherm was allowed to take the temperature to 195°C. The temperature was allowed to drop to 170°C and reaction then cooled to 130°C.
- 15 2. bb and bc were premixed and added. Reaction was held at 105°C over 1 hour, after which 51g of organic solvent was distilled off to remove the residual DEEA.
- 20 3. cc was then added and mixed in well over 30 mins. at 70°C.
4. dd was then slowly added while maintaining constant stirring.

The resulting coating composition exhibited excellent stability and had a viscosity of 31 sec. Ford Cup No. 4 at 42% solids at 25°C.

EXAMPLE 11

PREPARATION OF AMINE TERMINATED POLYETHER
USING 2-DIMETHYLAMINO-2-METHYLOL PROPANOL (DMAMP)

5 The method of preparation was the same as for
EXAMPLE 10 except that ingredient bb was DMAMP.

The resulting coating composition had excellent
stability and a viscosity of 23 sec. Ford Cup No. 4 at
37% solids and 25°C.

EXAMPLE 12

10 PREPARATION OF AMINE TERMINATED POLYETHER
USING DIMETHYL ISOPROPANOLAMINE (DMPA)

The method of preparation was the same as for
EXAMPLE 10 except that ingredient bb was DMPA.

15 The resulting coating composition had excellent
stability and a viscosity of 24 sec. Ford No. 4 Cup at
34% solids at 25°C.

EXAMPLE 13

AQUEOUS DISPERSION BASED ON TRIETHANOLAMINE
TERMINATED POLYETHER

20 The method of preparation was the same as for
EXAMPLE 10 except that ingredient bb was
triethanolamine.

The resulting coating composition had excellent
stability and a viscosity of 32 sec. Ford No. 4 Cup at
25 34.8% solids at 25°C.

The following Examples demonstrate that
modified epoxy resins with a wide range of epoxy

equivalent weights, can be successfully amine terminated and dispersed in water using the invention herein disclosed.

EXAMPLES 14 TO 20

5 PART 1: MODIFICATION OF THE EPOXY RESIN

In each of these Examples, the appropriate mole ratio (see TABLE 1) of Bisphenol A and DER343, to a total weight of 803g, were mixed with 162g of butyl cellosolve, in a 2-litre flange-topped, round-bottomed flask, fitted with a condenser, a stirrer and a temperature probe which was connected to a temperature controller. The components were mixed continuously while heating to 170°C over 1 hour, after which the heating was turned off and the exotherm allowed to take the temperature to reflux (195 - 200°C). The reaction was allowed to cool to 100°C and samples were taken for epoxy equivalent weight determination.

PART 2: AMINE TERMINATION

After the samples were taken a mixture of DMEA and butyl cellosolve was added to the vessel and the temperature maintained at 100°C for 1 hour. The mixtures used were as follows:

	<u>DMEA (g)</u>	<u>Butyl Cellosolve (g)</u>
EXAMPLE 10	190	98
25 EXAMPLE 11	190	49
EXAMPLE 12	98	98
EXAMPLES 14 TO 19	49	49

The reaction mixture was then distilled for 1 hour under a stream of nitrogen to remove excess DMEA.

PART 3: DISPERSION

5 200g of the product of Part 2 was in each case mixed thoroughly with 178g of carboxylic acid functional polymer for 1 hour at a temperature of 70°C. Then in each case water, 450g, was stirred into the reaction mixture and a uniform dispersion obtained.

10

TABLE 1

RESULTS ON EXAMPLES 10 TO 12 AND 14 TO 19

EXAMPLE NO.	10	11	12	14	15	16	17	18	19
Moles of Bisphenol A for each mole of DEFGH3	0.5	0.67	0.82	0.93	1	1.08	1.22	1.5	2.0
Epoxy equivalent wt. of modified epoxy resin	500	880	1900	3000	4000	6000	8000	10000	28000
Calculated molecular wt. of modified epoxy resin	1000	1600	3000	5000	4000	3500	2000	1400	1000
Viscosity of final aqueous dispersion sec. Ford Cup No.4 at 25°C	>300	88	119	26	15	20	19	18	18
Measured solids	32.9	32.8	32.3	32.9	33.1	34.1	34.3	34.3	33.5

EXAMPLE 20

PREPARATION OF AMINE TERMINATED POLYETHER
FROM MODIFIED EPOXY RESIN HAVING VERY
HIGH EPOXIDE EQUIVALENT WEIGHT

			<u>Wt. (g)</u>
5	a a	DER343	1500
	a b	Bisphenol A	1289
	a c	Butyl cellosolve	782
	b b	DMEA	148
10	b c	Butyl cellosolve	148
	c c	Carboxylic acid functional polymer (Example 5)	3390
	d d	Water	6704

1. aa, ab and ac were mixed in a flask fitted with a condenser, a thermometer-temperature controller, and a stirrer. The mixture was heated to 170°C, and the heating turned off. The exotherm was allowed to take the temperature up to 190°C (reflux temp.), after which the reaction was kept at reflux temperature for 5 hours. After this time the modified epoxy resin was found to have epoxy equivalent weight of 65000. The mixture was then cooled to 130°C.
2. bb and bc were premixed, then added into the flask; reaction was maintained at 105°C over 1 hour, after which 148g of organic solvent was distilled off.

3. cc was then added and mixed in well. The above resin blend was dispersed into water by adding dd slowly with constant stirring. The viscosity of the paint was 126 sec. Ford Cup No. 4 at 37% solids at 25°C.

5

EXAMPLE 21

DISPERSION OF TITANIUM DIOXIDE INTO COATING

		<u>Wt.</u>
	a a Coating composition of Example 10	1026
	a b Titanium Dioxide	214
10	b b Butanol	20
	c c Water	20

1. aa and ab were ground for 30 minutes in a speed mill and a grind of less than 6 microns obtained.
2. bb was added to eliminate foam and cc to adjust viscosity.
- 15

The pigmented coating composition had good stability and a viscosity of 28 sec. Ford No.4 Cup at 50% solids and 25°C.

EXAMPLE 22

DISPERSION OF A POLYMER BLEND CONTAINING
A HIGH LEVEL OF AMINE TERMINATED POLYETHER

			<u>Wt. (g)</u>
5	a a	Amine terminated polyether (Example 7)	213.2
	b b	Carboxylic acid functional polymer	9.20
	c c	DMEA	0.88
10	d d	Water	154

1. The carboxylic acid functional polymer used was prepared as follows. 840g acrylic acid, 767g styrene, 11g benzoyl peroxide and 3055g butyl cellosolve were added to a 5-litre
15 five-necked, flange-topped, round-bottomed flask fitted with a condenser, a stirrer and a thermometer, stirred until ac had dissolved, and heated to 80°C. The heating was then turned off, the exotherm was allowed to take
20 the temperature to reflux after which the temperature started to drop. When the temperature had dropped to 130°C, the heating was turned on and 273g solvent distilled off. The resulting polyelectrolyte had an NVM of 35%. When a sample
25 was diluted with an equal weight of butyl cellosolve, it had a Gardner-Holdt viscosity of W at 25°C.
2. aa and bb were mixed well in a 750 ml flask at 70°C.

3. cc was added with constant stirring.
4. dd was added over 30 mins.

The dispersion was stable and had a viscosity of 180 sec. Ford Cup No.4 at 25°C at 46% solids.

CLAIMS

1. An aqueous dispersion of a tertiary amine terminated polyether and a carboxylic acid functional polymer wherein the tertiary amine terminated polyether is the product of the reaction of the epoxy group of an epoxy resin with the hydroxy group of a hydroxy functional tertiary amine and the carboxylic acid functional polymer is present in an amount sufficient to render the combination water dispersible.
2. An aqueous dispersion according to claim 1 in which the amine terminated polyether is formed from a Bisphenol A type epoxy resin.
3. An aqueous dispersion according to claim 2 in which the molecular weight of the Bisphenol A type epoxy resin is increased by further reaction with Bisphenol A.
4. An aqueous dispersion according to any of the preceding claims in which the epoxy equivalent weight of the epoxy resin is greater than 5000.
5. An aqueous dispersion according to any of the previous claims in which the hydroxy functional tertiary amine is a mono hydroxy tertiary amine.
6. An aqueous dispersion according to any of the previous claims in which the hydroxy functional tertiary amine is N,N-dimethylethanolamine (DMEA).
7. An aqueous dispersion according to claim 1 in which the hydroxy functional tertiary amine is triethanolamine.
8. An aqueous dispersion according to any of the previous claims in which the carboxylic acid functional polymer is a polymer containing acrylic and/or methacrylic acid and/or crotonic acid.

9. An aqueous dispersion according to any of the previous claims in which the carboxylic acid functional polymer is a copolymer containing hydrophobic monomers such as styrene, butyl acrylate, ethyl acrylate, methyl methacrylate and butyl methacrylate.
10. An aqueous dispersion according to any of the previous claims in which the carboxylic acid functional polymer is a copolymer containing hydrophilic monomers such as hydroxy ethyl acrylate and hydroxy propyl methacrylate.
11. An aqueous dispersion according to any of the previous claims in which the carboxylic acid functional polymer contains acrylamide.
12. An aqueous dispersion according to any of the previous claims in which the carboxylic acid functional polymer is polymerised separately from the amine terminated polyether, and later mixed with said polyether.
13. An aqueous dispersion according to any of the previous claims in which the carboxylic acid functional polymer is polymerised in the presence of the amine terminated polyether.
14. A process for producing an aqueous dispersion of a tertiary amine terminated polyether and a carboxylic acid functional polymer which comprises:
 - a) reacting the epoxide groups of an epoxy resin with a hydroxy functional tertiary amine to form a tertiary amine terminated polyether, and
 - b) dispersing the tertiary amine terminated polyether in water with the aid of a carboxylic acid functional polymer in an amount sufficient to render the combination dispersible.
15. A process as claimed in claim 14, wherein the tertiary amine terminated polyether is formed under substantially anhydrous reaction conditions.

16. A process as claimed in claim 14, wherein the carboxylic acid functional polymer is dissolved in water prior to being used to disperse the tertiary amine terminated polyether.

17. A coating composition comprising the aqueous dispersion of any of claims 1 to 13.

18. A coating composition comprising a stable dispersion in water of:

i) a tertiary amine terminated polyether which is in ionic association with

ii) a polymer derived, at least in part, from carboxylic acid functional monomers, and

iii) a crosslinking agent.

19. A coating composition as claimed in claim 18, formulated for use in the lining of metal containers for food and beverages.

CLAIMS

1. A process for producing an aqueous dispersion of a tertiary amine terminated polyether and a carboxylic acid functional polymer which comprises:
 - a) reacting the epoxide groups of an epoxy resin with a hydroxy functional tertiary amine to form a tertiary amine terminated polyether, and
 - b) dispersing the tertiary amine terminated polyether in water with the aid of a carboxylic acid functional polymer in an amount sufficient to render the combination dispersible.
2. A process according to claim 1 in which the amine terminated polyether is formed from a Bisphenol A type epoxy resin.
3. A process according to claim 2 in which the molecular weight of the Bisphenol A type epoxy resin is increased by further reaction with Bisphenol A.
4. A process according to any of the preceding claims in which the epoxy equivalent weight of the epoxy resin is greater than 5000.
5. A process according to any of the previous claims in which the hydroxy functional tertiary amine is a mono hydroxy tertiary amine.
6. A process according to any of the previous claims in which the hydroxy functional tertiary amine is N,N-dimethylethanolamine (DMEA).
7. A process according to claim 1 in which the hydroxy functional tertiary amine is triethanolamine.
8. A process according to any of the previous claims in which the carboxylic acid functional polymer is a polymer containing acrylic and/or methacrylic acid and/or crotonic acid.

9. A process according to any of the previous claims in which the carboxylic acid functional polymer is a copolymer containing hydrophobic monomers such as styrene, butyl acrylate, ethyl acrylate, methyl methacrylate and butyl methacrylate.
10. A process according to any of the previous claims in which the carboxylic acid functional polymer is a copolymer containing hydrophilic monomers such as hydroxy ethyl acrylate and hydroxy propyl methacrylate.
11. A process according to any of the previous claims in which the carboxylic acid functional polymer contains acrylamide.
12. A process according to any of the previous claims in which the carboxylic acid functional polymer is polymerised separately from the amine terminated polyether, and later mixed with said polyether.
13. A process according to any of the previous claims in which the carboxylic acid functional polymer is polymerised in the presence of the amine terminated polyether.
14. A process according to any of the previous claims, wherein the tertiary amine terminated polyether is formed under substantially anhydrous reaction conditions.
15. A process according to any of the previous claims, wherein the carboxylic acid functional polymer is dissolved in water prior to being used to disperse the tertiary amine terminated polyether.

16. A process for preparing a coating composition which comprises forming a stable dispersion in water of:
- i) a tertiary amine terminated polyether which is in ionic association with
 - ii) a polymer derived, at least in part, from carboxylic acid functional monomers, and
 - iii) a crosslinking agent.

(19)



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(54) Water reducible coating compositions.

(57) An aqueous dispersion of a tertiary amine terminated polyether and a carboxylic acid functional polymer. The tertiary amine terminated polyether is the product of the reaction of the epoxy group of an epoxy resin with the hydroxy group of a hydroxy functional tertiary amine and the carboxylic acid functional polymer is present in an amount sufficient to render the combination water dispersible.

A process for producing the aqueous dispersion is also described, together with coating compositions based on the dispersion. These coating compositions are particularly useful in the lining of metal containers for food and beverages.

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EUROPEAN SEARCH REPORT

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DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.4)
Y	GB-A-2 087 895 (THE INTERNATIONAL PAINT COMPANY LTD) * Claims; page 1, lines 98-120; page 2, lines 24-43; page 3, line 125 - page 4, line 18 *	1-12, 15, 18	C 09 D 5/02 C 09 D 3/58 C 08 G 59/64 C 08 L 63/00 C 08 J 3/02
Y-12	US-A-4 339 369 (D.D. HICKS et al.) * Claims *	1-12, 15, 18	
A	DE-A-2 033 770 (PPG INDUSTRIES) * Claims; page 3, paragraph 3 - page 4, paragraph 2 *	1-12	
A	DE-A-1 644 783 (THE DOW CHEMICAL CO) * Claims; page 5, paragraph 2 *	1-12	
			TECHNICAL FIELDS SEARCHED (Int. Cl.4)
			C 08 G C 09 D C 08 J
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 29-02-1988	Examiner VOIGTLAENDER R.O.J.
CATEGORY OF CITED DOCUMENTS			
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